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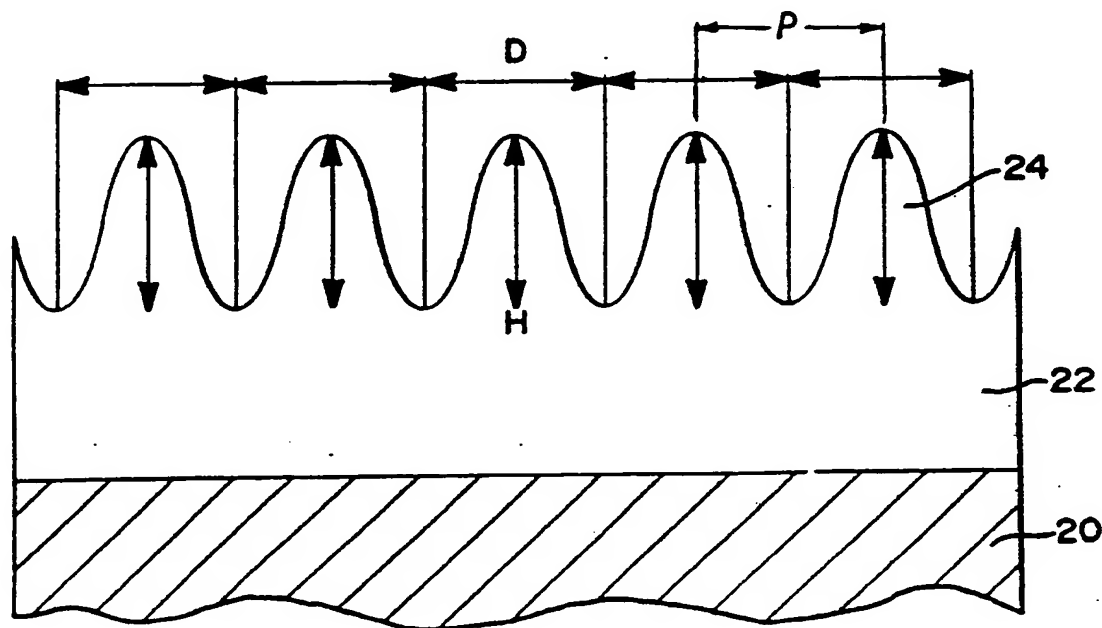
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<p>(21) International Application Number: PCT/US88/01675 (22) International Filing Date: 20 May 1988 (20.05.88) (31) Priority Application Number: 052,991 (32) Priority Date: 22 May 1987 (22.05.87) (33) Priority Country: US (71) Applicant: GLASSTECH SOLAR, INC. [US/US]; 12441 West 49th Avenue, Wheat Ridge, CO 80033 (US). (72) Inventors: YABA, Susumu ; 543, Sanmaicho, Kanagawa-ku, Yokohama-shi, Kanagawa-Pref. (JP). WALKER, Christopher, M. ; 856 18th Street, Boulder, CO 80302 (US). MUHL, Stephen ; 1237 C.S. Flower Circle, Lakewood, CO 80226 (US). MADAN, Arun ; 327 Lamb Lane, Golden, CO 80401 (US).</p>		<p>(74) Agent: LAFONTAINE, Earl, J.; Brooks & Kushman, 2000 Town Center, Suite 2000, Southfield, MI 48075 (US). (81) Designated States: AT (European patent), BE (European patent), CH (European patent), DE (European patent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent). Published <i>With international search report.</i></p>

(54) Title: SOLAR CELL SUBSTRATE



(57) Abstract

A solar cell substrate comprising a glass substrate (20) and a transparent electrically conductive layer (22) formed thereon, said conductive layer having a plurality of polygonal projections (24), having approximate diameters of from 0.1 to 0.3 μm and height/diameter ratios of at least 0.6.

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SOLAR CELL SUBSTRATETechnical Field

The present invention relates to a solar cell substrate useful in the production of amorphous silicon solar cells.

Background Art

As shown in Figure 1, a prior art amorphous silicon solar cell is illustrated generally as 8 and comprises a transparent insulating substrate 1, a transparent electrically conductive layer 6 composed of a p-type amorphous silicon layer 3, an i-type amorphous silicon layer 4, an n-type amorphous silicon layer 5, and an aluminum electrode 7 acting as the back contact. This configuration is practically used as a photoelectric conversion device capable of being produced at a relatively low cost. Such an amorphous solar cell 8 is designed so that light enters the solar cell through the transparent insulating substrate 1 and is absorbed mainly by the i-type amorphous silicon layer 4. An electromotive force is generated between the two electrodes, the transparent electrically conductive layer 2 and the aluminum electrode 7, and electricity is led out of the solar cell by a conductor 10.

For such an amorphous silicon solar cell, it is most important to improve the photoelectric conversion efficiency. As a method of achieving this, it is known to roughen the surface of the transparent electrically conductive layer. By roughening the surface, incident light is scattered at the interface between the transparent electrical-

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ly conductive layer and the amorphous silicon semiconductor layer. The effect of this optical scattering is to reduce the surface reflection loss of the incident light and to increase the absorbance in the i-type amorphous silicon layer. As a result of multiple reflection and refraction in the amorphous silicon semiconductor layer, the length of the optical path increases and the light-collecting efficiency of the amorphous silicon solar cell for long wavelength lights will be improved by the light-trapping effect in the i-type amorphous silicon layer. The short-circuit current density will also be increased, thereby improving the power generation efficiency and the photoelectric conversion efficiency.

As specific methods for such surface roughening, Japanese Unexamined Patent Publication No. 57756/1983 discloses an average particle size of the surface of the transparent electrically conductive layer from 0.1 to 22.5 μ m; Japanese Unexamined Patent Publication No. 201470/1984 discloses the preparation of a transparent electrically conductive layer composed of crystal grains having an average grain size of from 0.1 to 0.9 μ m; Japanese Unexamined Patent Publication No. 103384/1984 discloses an average grain size of fine crystals constituting a transparent electrically conductive layer to be at least 300 Å; and Japanese Unexamined Patent Publication No. 175465/1985 discloses forming a silicon oxide coating layer having a haze of at least 1% and having a number of projections having a diameter of from 300 to 500 Å and a height of from 200 to 3,000 Å, to obtain a solar cell substrate having a transparent electrically conductive layer.

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With solar cell substrates having the above-mentioned structures, it may be possible to increase the photoelectric conversion efficiency to some extent, but they are inadequate for the purpose of greatly improving the efficiency.

Summary of Invention

It is an object of the present invention to provide a solar cell substrate coated with a transparent conductive oxide having an optimum textured surface whereby an improved photoelectric conversion efficiency can be obtained.

The present invention provides a solar cell substrate including a glass substrate and a transparent electrically conductive layer formed thereon. The transparent electrically conductive layer has a plurality of polygonal projections having diameters of from 0.1 to 0.3 μ m and height/diameter ratios of at least 0.6.

Brief Description of Drawings

Figure 1 is a cross-sectional view of an amorphous silicon solar cell; and

Figure 2 is a partial enlarged cross-sectional view illustrating the surface of the transparent electrically conductive layer of the solar cell substrate of the present invention.

Best Mode for Carrying Out The Invention

For the substrate to be used in the present invention, it is possible to employ a transparent glass sheet made of soda-lime silicate glass, aluminosilicate glass, borate glass, lithium aluminosilicate glass, quartz glass or various other glasses. It is desirable for the substrate to

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exhibit a high transmittance, e.g. transmittance of at least 80%, in a wavelength range of from 350 to 800 nm. It is also desirable to have good insulating properties, high chemical and physical durability and excellent optical properties. In the case of a substrate made of a glass containing sodium, such as soda-lime silicate glass, or a substrate made of a low alkali-containing glass, an alkali barrier coating, such as a silicon oxide, aluminum oxide or zirconium oxide layer, may be provided on the substrate in order to minimize diffusion of ions, including sodium, from the glass into the transparent electrically conductive layer formed thereon.

The transparent electrically conductive material formed on the glass substrate is preferably made of a transparent metal oxide having a desired conductivity, such as tin oxide doped with from 0.1 to 5% by weight of fluorine, tin oxide doped with from 0.1 to 30% by weight of antimony, or indium oxide doped with from 0.5 to 30% by weight of tin. Among those described, a transparent electrically conductive layer made of fluorine-doped tin oxide is most suitable for a solar cell substrate, since it is possible to readily obtain a sheet resistance lower than $30 \Omega / \square$. The fluorine-doped tin oxide is resistant to chemical reduction caused by hydrogen plasma, to which it is exposed during the formation of amorphous silicon layers in a plasma enhanced chemical vapor deposition chamber.

Referring now to Figure 2, the solar cell substrate shown has a transparent electrically conductive layer with a roughened texture over its entire surface. Namely, the transparent electrically conductive layer 22 has a plurality of projec-

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tions 24, each having a diameter D of from 0.1 to 0.3 μm , and ratio of a height H to a diameter D of at least 0.6, preferably from 0.7 to 1.2. The height H of such projections is preferably from 0.1 to 0.6 μm , more preferably less than 0.25 μm . Further, the projections 24 are preferably formed with a pitch P (summit to summit distance) of from 0.1 to 0.5 μm , for, if the projections are too far apart from each other, the transparent electrically conductive layer does not effectively cause light scattering. The shape of the projections vary. In a typical example, the shape of the base portion of each projection is tetragonal or has a shape of a higher order polygon. Atmospheric chemical vapor deposition automatically produces a textured oxide layer. The shape of the textured projections depend on the chemical composition of the layer.

In the present invention, projections having a diameter outside the range of from 0.1 to 0.5 μm or a height/diameter ratio outside the range of at least 0.6, may be present to some extent so long as more than 80% of the surface topography conforms to the preferred specifications.

The transparent electrically conductive layer should have a sheet resistance value less than $30\Omega/\square$, preferably from 4 to $10\Omega/\square$. This range is desirable particularly in view of avoiding ohmic losses in a large area substrate. In the present invention, the haze is most preferably from 8 to 30% in view of the above-mentioned relationship between the particle shape and the optical properties needed to enhance the photoelectric conversion efficiency of the solar cells.

The transparent electrically conductive layer of the present invention may be prepared by

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various conventional methods, such as chemical vapor deposition, pyrolytic spraying, sputtering, vacuum deposition, ion plating or dipping. Of these methods, chemical vapor deposition and sputtering are most preferred, since projections having the above-mentioned diameter, height and height-diameter ratio, can readily be obtained in a controlled manner.

The invention is illustrated by the following examples, but is not intended to be limited to the details therein.

Examples

(1) Preparation of SnO_2 layer

Nine samples of solar cell substrates differing from one another in the surface texture of the transparent electrically conductive layer were fabricated.

Each substrate consisted of a soda-lime silicate glass sheet (4 inch x 3.5 inch x 1.1 mm thick) and a SiO_2 layer of 800 Å thick formed thereon.

Preparation of transparent electrically conductive layer of SnO_2 was accomplished by using SnCl_4 , H_2 , CH_3OH , and HF carried by nitrogen gas, as reactive gases.

The SnO_2 layer was prepared in a belt conveyor atmospheric chemical vapor deposition furnace having both heating and cooling zones. The heating zone included a four inch wide nozzle for supplying the reactive gases. This nozzle has five fine slit gas outlets; SnCl_4 , and dilution nitrogen flows out of slit number three; the middle slit; H_2O , CH_3OH , HF and their dilution nitrogen flows out of slits 1 and 5; nitrogen to separate the groups of

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reactants flows out of slits 2 and 4. Furthermore, the distance between the glass substrate and the gas outlets is preferably adjustable.

Sample No. 1

5 A glass substrate coated with a SiO_2 layer
to a thickness of 800 Å, was introduced into the
above CVD furnace at a belt conveyor speed of 0.20
m/min. and heated to a substrate temperature of 600
°C. The reaction gases were then blown onto the
10 substrate by the gas supply nozzle to form the SnO_2
layer. The gas compositions were 0.025 liter/min.
of SnCl_4 and 1.5 liter/min of N_2 from the central
slit; 1.5 liter/min. of N_2 from the separating
lines; and 0.93 liter/min. of H_2O , 0.015 liter/min.
15 of HF , 0.017 liter/min. of CH_3OH and 1.5 liter/min.
of N_2 from the outerlying slits.

Sample 2 to Sample 9 were fabricated in
the same CVD furnace with the conditions shown in
Table 1.

20 (2) Measurement of the surface texture

(i) Height of projections:

 The total thickness, t_1 , of the trans-
parent electrically conductive layer formed on the
substrate was initially measured with a feeler-type
25 thickness meter (Taly-step; Taylor-Hobson Co.,
needle tip 12.5 μm in diameter). Then an area of
the layer was abraded with a 0.5 μm diamond paste
under a load of 500 g/cm². The layer thickness and
the haze in this area were measured at intervals
30 during the abrasion operation, and the layer
thickness t_1 at the time when the haze became 2% or
less was determined. The difference between the

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initial thickness t_0 and the thickness t_1 , i.e. $t_1 - t_0$, is equal to the height of the projections.

(ii) Diameter of projections:

5 A scanning electron micrograph of the transparent electrically conductive layer formed on the substrate was taken, and the diameter of the projections was measured from the micrograph.

(iii) Table 2 shows the measured values of H, D and H/D for each sample.

10 The photoelectric conversion efficiency, the short-circuit density, the open-circuit voltage, and the fill factor of solar cells made on sample are also shown.

15 Samples 1 to 7 all have H, D and H/D ratios within the range of the present invention. Samples 8 and 9 have H, D or H/D ratios outside the range of the present inventions and it can be seen that the properties of the solar cells made on these samples are inferior to those of the solar cells made on the samples 1 through 7.

20 The solar cells consisted of the transparent electrically conductive layer of each sample, a p-type amorphous silicon layer (thickness: about 100 Å), an i-type amorphous silicon layer (thickness: about 4,000 Å), a n-type amorphous silicon layer (thickness: about 1,000 Å), and a metal electrode layer (thickness: about 10,000 Å).

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Table 1. Condition of SnO_2 layer deposition

Sample No.	Substrate Temperature (°C)	Belt Conveyor Speed (m/min.)	Central Stream		Separating Stream N_2 (l/min.)	Outer stream			
			SnCl_4 (l/min.)	N_2 (l/min.)		H_2O (l/min.)	HF (l/min.)	CH_3OH (l/min.)	N_2 (l/min.)
1	600	0.20	0.025	1.6	1.5	0.93	0.015	0.017	1.5
2	585	↑	↑	↑	↑	↑	0.005	0.007	↑
3	↑	↑	↑	↑	↑	↑	0.015	0.017	↑
4	615	0.11	↑	↑	↑	↑	↑	↑	↑
5	540	0.20	↑	↑	↑	↑	↑	↑	↑
6	585	↑	↑	↑	↑	↑	0.005	↑	↑
7	600	↑	↑	↑	↑	1.26	0.015	0.035	↑
8 (comparative example)	550	↑	↑	↑	↑	0.48	↑	0.017	↑
9 (comparative example)	615	0.11	↑	↑	↑	0.93	0.030	↑	↑

Table 2. Surface Texture and Solar Cell Property

Sample No.	Surface properties of transparent electrically conductive layer (average values)			Sheet resistance (specific resistance) of transparent electrically conductive layer (Ω/\square)	Mass of transparent electrically conductive layer (g)	Thickness and visible light transmittance of transparent electrically conductive layer	Solar cell properties			
	Height of projections (μ)	Diameter of projections (μ)	Ratio of height to diameter				Photoelectric conversion efficiency η (%)	Short-circuit density (mA/cm^2)	Open-circuit voltage (V)	Fill factor (%)
1	1400	1500	0.93	8.2	13.6	5820 Å 79.0%	9.8	16.5	0.815	0.73
2	2900	2600	1.12	5.2	25.0	8500 Å 82.0%	9.4	16.4	0.81	0.71
3	1000	1400	0.71	8.0	9.7	5620 Å 80.5%	9.36	16.48	0.797	0.70
4	2430	3000	0.81	6.3	36.0	12200 Å 76.0%	9.2	16.1	0.79	0.72
5	740	1000	0.74	9.9	4.8	5180 Å 80.9%	9.1	16.0	0.82	0.70
6	1400	1000	1.40	7.1	21.0	5900 Å 80.2%	9.0	16.0	0.80	0.70
7	1500	2500	0.60	8.0	14.0	8100 Å 78.0%	8.9	15.2	0.84	0.69
8 (comparative Example)	800	1500	0.53	8.7	5.6	4800 Å 83.6%	8.53	15.55	0.813	0.67
9 (comparative Example)	4030	3100	1.30	4.7	63.5	10700 Å 76.7%	8.0	14.8	0.81	0.67

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In the present invention, the incident light enters the solar cell through the interface of the transparent electrically conductive layer and the amorphous silicon layer and penetrates into the amorphous silicon layer. When the electrically conductive layer has a textured surface, the light is scattered by the textured surface and the scattered light is taken into the amorphous layer. The optical path in the silicon layer is elongated, and the absorption efficiency of light is thereby improved. This effect is particularly remarkable in a long wavelength range where the absorptive coefficient of silicon is small. In order to increase the scattering of light in this long wavelength range, certain levels of the height and the diameter of projections are required.

The transparent conductive layer of the present invention has a plurality of projections having diameters of from 0.1 to 0.3 μ m and height/diameter ratios of at least 0.6, preferably from 0.7 to 1.2.

It is desirable for the diameter and height/diameter ratio to be quite large since if the diameter is less than 0.1 μ m and/or the height/diameter ratio is less than 0.6, the projections become too small and/or too flat to efficiently produce light scattering.

However, if the diameter is larger than 0.3 μ m, and/or the height/diameter ratio is larger than 1.2, the height becomes too great, which makes it difficult to uniformly deposit an amorphous silicon layer on the transparent electrically conductive layer. In this respect, diameter and height/diameter ratio are restricted by the thickness of the amorphous silicon layer or the

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ability of the amorphous silicon deposition system to conformally deposit layers while maintaining good electrical properties.

5 Similarly, to form projections having diameters of larger than $0.3\mu\text{m}$, it is necessary to heat the glass substrate up to high temperatures, which is not preferable for a glass substrate.

10 Comparing the sample 1-7 of the present invention with the comparative samples, 8, 9, it is easily seen that the performance of the solar cells prepared on the substrate of the present invention provide the following advantages:

- 15 (1) the short circuit current density is improved by about 1 mA/cm^2 ; and
(2) the photoelectric conversion efficiency is improved by about 0.5% absolute units.

20 While the best modes for carrying out the invention have herein been described in detail, those familiar with the art to which this invention relates will recognize various alternatives and embodiments for carrying out the invention as defined by the following claims.

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WHAT IS CLAIMED IS:

1. A solar cell substrate, comprising a glass substrate and a transparent electrically conductive layer formed thereon, said conductive layer having a plurality of projections, having diameters of from 0.1 to 0.3 μ m and height/diameter ratios of at least 0.6.
2. The solar cell substrate as in claim 1, wherein the height/diameter ratio of the projections is from 0.7 to 1.2.
3. The solar cell substrate as in claim 1, wherein the transparent electrically conductive layer is selected from the group consisting of tin oxide, fluorine-doped tin oxide, antimony-doped tin oxide and indium tin oxide.
4. The solar cell substrate as in claim 1, wherein the transparent electrically conductive layer has a sheet resistance of from 4 to 10 Ω / \square .
5. The solar cell substrate as in claim 1, wherein the transparent electrically conductive layer has a haze as from 8 to 30%.

FIG 1
PRIOR
ART

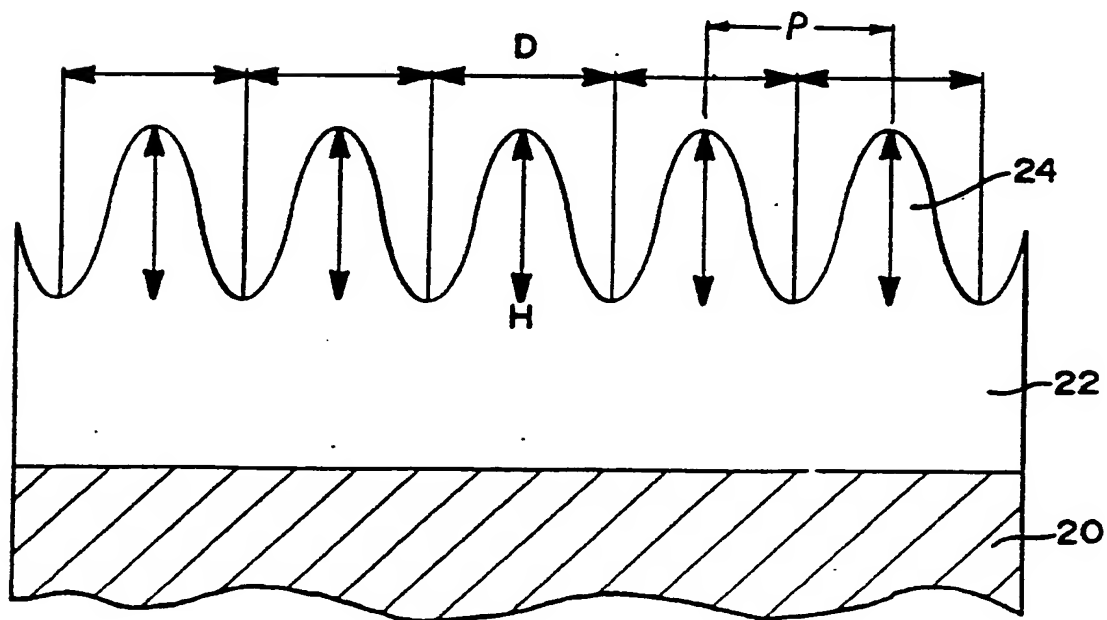
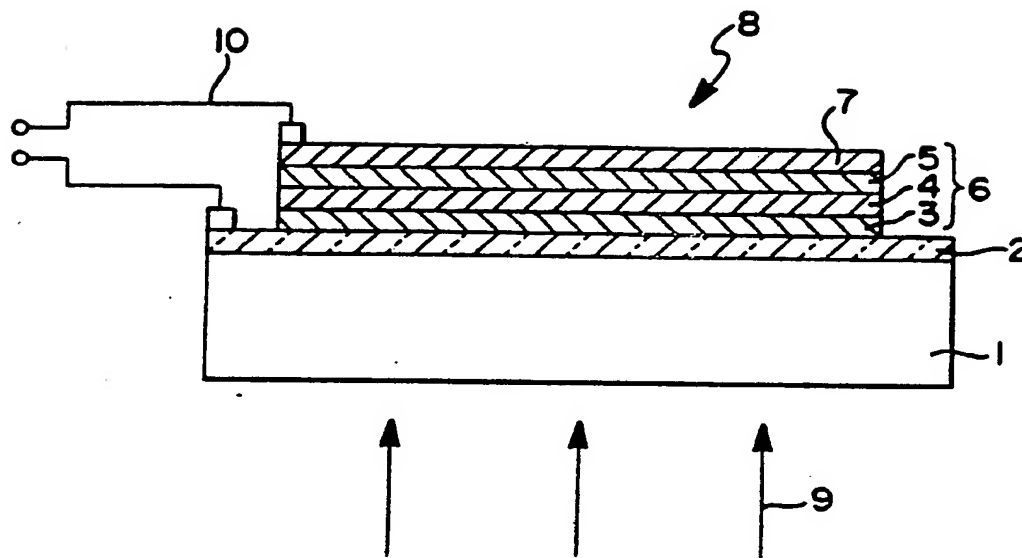


FIG 2

INTERNATIONAL SEARCH REPORT

International Application No PCT/US88/01675

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ³		
According to International Patent Classification (IPC) or to both National Classification and IPC		
INT. CL ⁴ B32B 17/06; H01L 31/04		
U.S. CL 428/432; 136/256		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁴		
Classification System	Classification Symbols	
US	428/141; 174; 432 136/256; 258AM	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁴		
III. DOCUMENTS CONSIDERED TO BE RELEVANT ¹⁴		
Category ⁵	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
X	US, A 4,532,537 Issued 30 July 1985, (KANE) (see col. 2, line 65 through col. 3, line 16).	1-5
X	US, A 4,554,727 Issued 26 November 1985, (DECKMAN ET AL) (see fig. 1, col. 6, lines 33-42 and col. 14, lines 44-68).	1-5
X	US, A 4,599,482 Issued 08 July 1986, (YAMAZAKI) (see figs. 1-3 and col. 6, lines 26-52).	1-5
X,P	US, A 4,689,438 Issued 25 August 1987, (FUKATSU ET AL) (see fig. 1 and col. 3, lines 3-48).	1-5
X	A, US 4,500,743 Issued 19 February 1985, (HAYASHI ET AL) (see entire patent).	1-5
A	N, IEEE Electron Device Letters, vol. EDL-3, issued May 1982, H. Iida et al "High Efficiency a-Si: H p-i-n Solar Cell Using a SnO ₂ /Glass Substrate, pages 114-115.	
<p>⁵ Special categories of cited documents: ¹⁵</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"A" document member of the same patent family</p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search ²		Date of Mailing of this International Search Report ²
06 July 1988		22 AUG 1988
International Searching Authority ¹		Signature of Authorized Officer ²⁰
ISA/US		Aaron Weisstuch

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category*	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No
X	N, IEEE Electron Device Letters, vol. EDL-4, issued May 1982, H. Iida et al, "Efficiency of the a-Si:H Solar Cell and Grain Size of SnO ₂ Transparent Conductive Film", pages 157-159 (see entire article).	1-5
X	N, IEEE Transactions on Electron Devices, vol. ED-31, issued May 1984, H. Iida et al, "Optical Confinement Type a-Si:H Solar Cell Using Milky Tin Oxide on Glass", pages 717-719 (see entire article).	1-5
X	N, IEEE Transactions On Electron Devices, vol. ED-34, issued February 1987, H. Iida et al, "The Structure of Natively Textured SnO ₂ Film and its Application to an Optical Confinement-Type a-Si:H Solar Cell", pages 271-276 (see entire article).	1-5
X,P	US, A 4,694,116 Issued 15 September 1987, HAYASHI ET AL (see entire patent).	1-5
X,P	US, A 4,732,621 Issued 22 March 1988, MURATA ET AL (see fig. 8, cols. 1, 3-4).	1-5
A,E	US, A 4,746,372 Issued 24 May 1988, TAJIKA ET AL (see entire patent).	1-5